



UNIVERSITY OF PETROLEUM AND ENERGY
STUDIES

End Semester Examination, December 2021

Program: MSc Chemistry

Semester: III

Course: General Pharmacology

Duration: 03 hours

Course Code: HSPT8009

Max. Marks: 100

Instructions: All questions are compulsory

	SECTION A (Type the answers in test box)	(5Q x4M= 20 Marks)	CO
	MCQs, One or two line answers.		
Q1	A. Maximum first pass metabolism is seen by which route of administration a. Intravenous b. Intramuscular c. Intradermal d. Oral B. The parameters required to calculate renal clearance a. Rate of elimination and half-life b. Half-life and plasma concentration c. Half-life and volume of distribution d. Rate of elimination and plasma concentration C. Therapeutic index of the drug is an indicator of its- a. Potency b. Safety c. Toxicity d. Efficacy D. EC ₅₀ is the measure of a. Toxicity b. Potency c. Efficacy d. Bioavailability	4	CO1

Q2	<p>A. pKa is the value at which the drug is-</p> <ol style="list-style-type: none"> 50% ionized 10% ionized 90% ionized 25% ionized <p>B. Acidic drugs bind to</p> <ol style="list-style-type: none"> Globulin Alpha-1 glycoprotein Albumin None <p>C. A highly ionized drug in</p> <ol style="list-style-type: none"> Is excreted mainly by the kidney Can cross blood brain easily Is well absorbed from the intestine Accumulates in the cellular lipids <p>D. Most of the drug are excreted in-</p> <ol style="list-style-type: none"> Faeces Urine Saliva Sweat 	4	CO2
Q3	<p>A. Zero order kinetic is independent of-</p> <ol style="list-style-type: none"> Plasma concentration Clearance Volume of distribution Half-life <p>B. What is bioavailability?</p> <p>C. Define clearance.</p> <p>D. Define therapeutic index.</p>	4	CO2
Q4	<p>A. In which of the following clinical trial phases ethical clearance is not required-</p> <ol style="list-style-type: none"> Phase I 	4	CO3

	<p>b. Phase II</p> <p>c. Phase III</p> <p>d. Phase IV</p> <p>B. Define essential drugs.</p> <p>C. Give two examples of counterirritant used as drugs.</p> <p>D. What is teratogenicity?</p>		
Q5	<p>A. What is adverse drug reaction?</p> <p>B. What are prodrugs?</p> <p>C. Explain the term orphan drugs.</p> <p>D. Define pharmacovigilance.</p>	4	CO3
	<p>SECTION B</p> <p>(Scan and upload)</p>	<p>(4Qx10M=40 Marks)</p>	<p>CO</p>
Q1	<p>A. Write a note on sources of drugs</p> <p>B. Classify routes of drug administration</p>	5+5	CO1
Q2	<p>A. Elaborate factors effecting bioavailability</p> <p>B. What are the clinical significance of plasma protein bounding</p>	5+5	CO2
Q3	<p>Write a note on</p> <p>a. Penetration of drugs into brain and CSF</p> <p>b. Zero and first order kinetics</p>	5+5	CO3
Q4	<p>A. Describe receptor occupation theory</p> <p>B. Write on approaches to drug discovery/ invention</p>	5+5	CO4
	<p>SECTION C</p> <p>(Scan and upload)</p>	<p>(2Qx15M=30 Marks)</p>	<p>CO</p>
Q1	<p>What is biotransformation? Give details on phases in biotransformation and their subtypes.</p>	20	CO2
Q2	<p>Define receptors. Write a detailed note of transducer mechanism of drug action.</p>	20	CO3